## H. CON. RES. 299

## IN THE SENATE OF THE UNITED STATES

July 16, 2008

Received and referred to the Committee on Health, Education, Labor, and Pensions

## **CONCURRENT RESOLUTION**

Supporting the goals and ideals of National Cystic Fibrosis Awareness Month.

Whereas cystic fibrosis is one of the most common life-threatening genetic diseases in the United States and one for which there is no known cure;

Whereas the average life expectancy of an individual with cystic fibrosis is 37 years—an improvement relative to the 1960s when children with cystic fibrosis

- did not live long enough to attend elementary school, but still unacceptably short;
- Whereas approximately 30,000 people in the United States have cystic fibrosis, more than half of them children;
- Whereas one of every 3,500 babies born in the United States is born with cystic fibrosis;
- Whereas more than 10,000,000 Americans are unknowing, symptom-free carriers of the cystic fibrosis gene;
- Whereas the Centers for Disease Control and Prevention recommends that all States consider newborn screening for cystic fibrosis;
- Whereas the Cystic Fibrosis Foundation urges all States to implement newborn screening for cystic fibrosis to facilitate early diagnosis and treatment which improves health and life expectancy;
- Whereas prompt, aggressive treatment of the symptoms of cystic fibrosis can extend the lives of people who have the disease;
- Whereas recent advances in cystic fibrosis research have produced promising leads in gene, protein, and drug therapies beneficial to people who have the disease;
- Whereas innovative research is progressing faster and is being conducted more aggressively than ever before, due, in part, to the Cystic Fibrosis Foundation's establishment of a model clinical trials network;
- Whereas although the Cystic Fibrosis Foundation continues to fund a research pipeline for more than 30 potential therapies and funds a nationwide network of care centers that extend the length and quality of life for people with

cystic fibrosis, lives continue to be lost to this disease every day;

- Whereas education of the public about cystic fibrosis, including the symptoms of the disease, increases knowledge and understanding of cystic fibrosis and promotes early diagnosis; and
- Whereas the Cystic Fibrosis Foundation will conduct activities to honor National Cystic Fibrosis Awareness Month in May, 2008: Now, therefore, be it
- 1 Resolved by the House of Representatives (the Senate 2 concurring), That the Congress—
- 3 (1) honors the goals and ideals of National Cys-
- 4 tic Fibrosis Awareness Month;
- 5 (2) promotes further public awareness and un-6 derstanding of cystic fibrosis;
- (3) advocates for increased support for people
  with cystic fibrosis and their families;
- 9 (4) encourages early diagnosis and access to 10 high-quality care for people with cystic fibrosis to
- improve the quality of their lives; and

1	(5) supports	research to find a cure	for cystic
2	fibrosis by foster	ing enhanced research	programs
3	and expanded public-private partnerships.		
	Passed the House of Representatives July 15, 2008.		
	Attest:	LORRAINE C. MILLI	ER,
			Clerk.

By Robert F. Reeves,  $\label{eq:Deputy Clerk} Deputy \ Clerk.$